To: Egeghy, Peter[Egeghy.Peter@epa.gov]

From: NoReply@efsa.europa.eu Sent: Mon 4/27/2015 6:55:48 PM

Subject: Form Confirmation

Dear Peter Egeghy,

You requested access to the draft risk assessment report of the active substance "Glyphosate".

Please, find below the URL linking to the files of the rapporteur Member State assessment report.

Public consultation has not been started yet or is now closed on this active substance and no further comments can be accepted by EFSA or the respective rapporteur Member State.

Rapporteur Member State assessment report documents:

- Glyphosate RAR 01 Volume 1 2013-12-18 san.pdf
- Glyphosate RAR 02 Volume 2 2013-12-18 san.pdf
- Glyphosate RAR 03 Volume 3CA-CP B-1 2013-12-18 san.pdf
- Glyphosate RAR 04 Volume 3CA-CP B-2 2013-12-18.pdf
- Glyphosate RAR 05 Volume 3CA-CP B-3 2013-12-18.pdf
- Glyphosate RAR 06 Volume 3CA-CP B-4 2013-12-18.pdf
- Glyphosate RAR 07 Volume 3CA-CP B-5 2013-12-18 san.pdf
- Glyphosate RAR 08 Volume 3CA-CP B-6 2013-12-18 san.pdf
- Glyphosate RAR 09 Volume 3CA-CP B-7 2013-12-18.pdf
- Glyphosate RAR 10 Volume 3CA-CP B-8 2013-12-18.pdf
- Glyphosate RAR 11 Volume 3CA-CP B-8 Appendix 2013-12-18.pdf
- Glyphosate RAR 12 Volume 3CA-CP B-9 2013-12-18 san.pdf
- Glyphosate RAR 13 Volume 3CA-CP B-9 Appendix 2013-12-18.pdf
- Glyphosate RAR 39 LoEP CA-CP 2013-12-18.pdf
- Glyphosate RAR 40 list of studies version 1 CA-CP 2013-12-18.pdf

The documents regarding this substance are provided in compliance with the provisions of the relevant legislative framework 1.

EFSA would like to draw your attention to the preliminary nature of the risk assessment presented in the rapporteur Member State assessment report, as it represents the initial evaluation of the data by a rapporteur Member State which is subsequently peer reviewed by EFSA.

Yours sincerely,

The European Food Safety Authority

1 Existing active substances (2nd stage):

Art. 8(6) of the <u>Commission Regulation (EC) No 451/2000</u> as amended by <u>Commission Regulation (EC) No 1490/2002</u>

Existing active substances (3rd stage):

Art. 11(3) of <u>Commission Regulation (EC) No 1490/2002</u> as amended by <u>Commission Regulation (EC) No 1095/2007</u>

Existing active substances (4th stage):

Art. 24(5) of <u>Commission Regulation (EC) 2229/2004</u> as amended by <u>Commission Regulation (EC) No 1095/2007</u>

Resubmissions:

Art. 9 or Art. 19(2) of Commission Regulation (EC) No 33/2008

Annex I Renewals:

Art.11(2) of Commission Regulation (EC) No 737/2007